## **Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

- 1. (Currently amended) A composition for use in making commercial products, comprising Requol the R enantiomer of equal (R-equal).
- 2. (Original) The composition according to Claim 1 wherein the composition is made by isolating R-equol from a racemic mixture of S-equol and R-equol.
- 3. (Original) The composition according to Claim 1, consisting essentially of R-equol.
- 4. (Currently amended) The composition according to <u>Claim 1</u> [[Claim 3]] wherein the R-equol has an enantiomeric purity of 90% minimum enantiomeric excess (EE).
- 5. (Original) The composition according to Claim 4 wherein the R-equol has an enantiomeric purity of 96% minimum EE.
- 6. (Original) A food composition comprising an additive component comprising R-equol.
- 7. (Currently amended) [[A]] The food composition according to Claim 6, wherein the food comprises, per serving of food, at least about 1 mg, and up to about 300 mg, R-equol.
- 8. (canceled)
- 9. (Original) A composition for topical application to skin, comprising R-equol and a vehicle.
- 10. (Original) The composition for topical application to skin according to Claim 9, comprising by weight at least 0.1%, and up to 10%, of R-equol.

11. (Original) The composition according to Claim 9 where the R-equol is conjugated at the C-4'

or C-7 position to form a conjugate selected from the group consisting of glucuronide, sulfate,

acetate, propionate, glucoside, acetyl-glucoside, malonyl-glucoside, and mixtures thereof.

12. (Original) A method of delivering R-equol to a mammal to prevent or treat a disease or

associated condition, comprising administering to the mammal a composition comprising R-

equol or a conjugated analog thereof.

13. (Currently amended) The method according to Claim 12 where the composition is

administered in an amount sufficient to produce a transient level of R-equol [[S-equol]] in the

blood plasma of the mammal of at least 5 ng/mL.

14. (Original) The method according to Claim 12 where R-equol is conjugated at the C-4' or C-7

position to form a conjugate selected from the group consisting of glucuronide, sulfate, acetate,

propionate, glucoside, acetyl-glucoside, malonyl-glucoside, and mixtures thereof.

15. (Original) The method according to Claim 12 where the composition is administered to the

mammal orally in a dose amount of at least about 1 mg R-equol.

16. (Original) The method according to Claim 12 where disease comprises a hormone-dependent

disease or condition selected from group consisting of cardiovascular disease, diminished blood

vessel quality, lipid disorder, osteopenia, osteoporosis, liver disease, acute ovarian estrogen

deficiency, benign breast cancer, breast cancer, benign prostate cancer, prostate cancer, skin

cancer, colon cancer, vasomotor disturbances and night sweats associated with ovarian estrogen

deficiency or antiestrogen therapy, impaired cognition, dementia, and brain disorders manifest as

short or long-term memory loss.

17. (Original) The method according to Claim 16 wherein the hormone-dependent disease or

condition is selected from group consisting of cardiovascular disease, diminished blood vessel

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quality, lipid disorder, osteopenia, osteoporosis, liver disease, and acute ovarian estrogen

deficiency.

18. (Original) The method according to Claim 17 wherein the composition is administered in an

amount sufficient to reduce the level of lipids in the blood or serum.

19. (Original) The method according to Claim 17 wherein the composition is administered in an

amount sufficient to reduce the surrogate markers of bone turnover or prevent bone loss as

measured by bone mineral density.

20. (Original) The method according to Claim 17 wherein the composition is administered in an

amount sufficient to increase bone formation.

21. (Original) The method according to Claim 17 wherein the composition is administered in an

amount sufficient to prevent osteoporosis and reduce bone fracture.

Claims 22-24 (canceled)

25. (Original) The method according to Claim 12 where disease comprises a non-hormone-

dependent disease or condition selected from group consisting of inflammatory conditions of the

gastrointestinal tract, the prostate, the breast, the skin and bone, and a condition associated with

adenomatous polyps and familial polyposis.

26. (canceled)

27. (Original) The method according to Claim 25 wherein the non-hormone-dependent disease or

condition is selected from group consisting of inflammatory conditions of the gastrointestinal

tract, the prostate, the breast, the skin and bone.

28. (Original) The method according to Claim 12 wherein the composition is administered as a

food or food additive.

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29. (New) The composition according to Claim 1 where the R-equol is conjugated at the C-4' or

C-7 position to form a conjugate selected from the group consisting of glucuronide, sulfate,

acetate, propionate, glucoside, acetyl-glucoside, malonyl-glucoside, and mixtures thereof.

30. (New) The composition according to Claim 9 wherein topical composition further comprises

an agent selected from the group consisting of antifungals, vitamins, anti-inflammatory agents,

antimicrobials, analgesics, nitric oxide synthase inhibitors, insect repellents, self-tanning agents,

surfactants, moisturizers, stabilizers, preservatives, antiseptics, thickeners, lubricants,

humectants, chelating agents, skin penetration enhancers, emollients, fragrances, and colorants

and combinations thereof.

31. (New) The composition according to Claim 30, wherein the commercial composition

comprises by weight up to 10% of R-equol.

32. (New) The composition according to Claim 31, wherein the commercial composition

comprises by weight at least 0.1% of R-equol.

33. (New) The composition according to Claim 30 where the R-equal is conjugated at the C-4'

or C-7 position to form a conjugate selected from the group consisting of glucuronide, sulfate,

acetate, propionate, glucoside, acetyl-glucoside, malonyl-glucoside, and mixtures thereof.

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